

ACCESSION NR: AFLO17638

\$/0190/64/006/002/0265/0268

AUTHORS: Alekseyeva, I. A.; Semerneva, G. A.; Samarina, L. A.; Bulatov, M. A.; Spasskiy; S. S.

TITIE: The synthesis, polymerisation and copolymerization of polyorganosiloxanes containing methacrylate groups. 2. Investigation of polymerization and copolymerization by the infrared absorption spectra method

SOURCE: Vy=sokomolekulyarny=ye soyedineniya, v. 6, no. 2, 1964, 265-268

TOPIC TACS: organosilicon compound, organosiloxane, polyorganosiloxane, methacrylate, styrene, copolymer with styrene, methacrylate polysiloxane polymer, double bond, saturation of double bond, infrared spectra, absorption band, absorption band optical density

ABSTRACT: Block polymerization of methacrylate polysiloxanes (containing from zero to nine of the Si(Ch₃)₂O groups) and their copolymerization with styrene (in a ratio of 1 Mol of styrene monomer per 1 Hol of polysiloxane unit) were investigated. The polymerization was conducted in the presence of 0.2% benzoyl peroxide in sealed ampules, in an atmosphere of nitrogen, for 6 hours at 70 and 100C and 12 hours at 120C, when it underwent complete solidification. The infrared spectra Card 1/8

ACCESSION MR: APLO17638

were taken by means of a IKS-14 registering spectrophotometer, the absorption band at 1634 cm⁻¹ having been selected as representing the $\text{CH}_2 = \mathbb{C}$ — double bonds which decrease in numbers during the reaction process. The other band was the one at 697 cm⁻¹, which represents the $\text{Si}(\text{CH}_3)_2$ groups, the number of which remains constant. As can be seen from Fig. 1 on the Enclosure, an increase in the number of methylsiloxene groups causes the optical density ratios to drop due to a decrease in the double bond content. It is suggested that the presence of unreacted double bonds is due to steric hindrances. The copolymerisation with styrene was found to proceed towards an almost complete saturation of the double bonds. Orig. art. has: 2 charts and 1 table.

ASSOCIATION: Institut khimii Ural'akogo filiala AN SSSR (Institute of Chemistry, Ural Division AN SSSR)

SUPHITTED: 03Dec62

DATE ACQ: 23March.

EMCT: 01

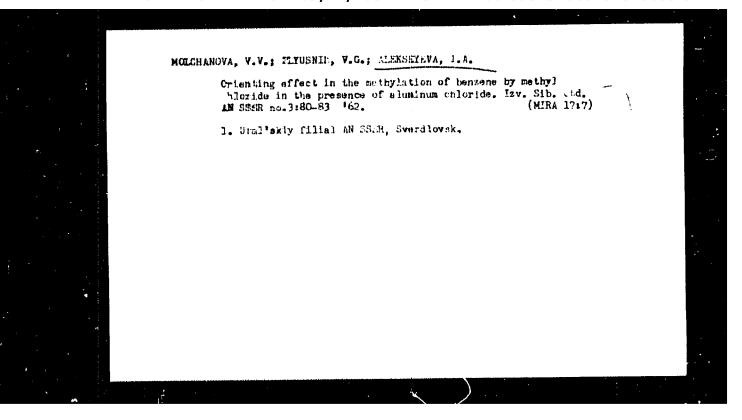
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OTHER: 002

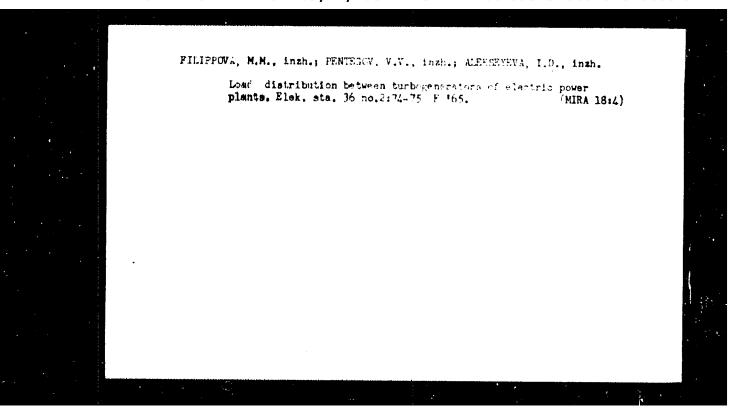
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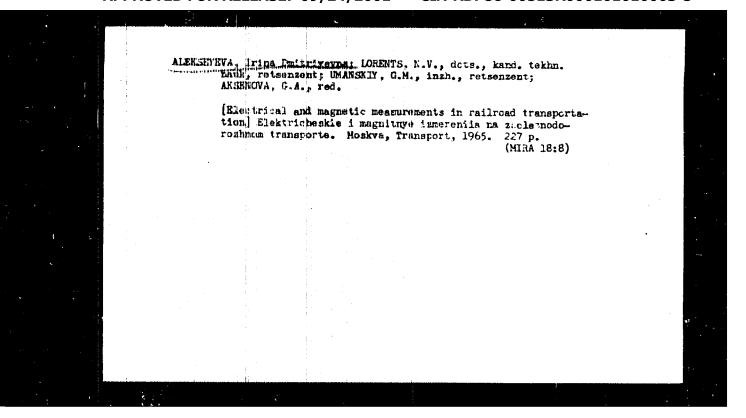


SEMENUEVA, G.A.; SUVOROV, A.L. SAMARINA, L.A.; ALEKSEYE/A, 1.A.; SPASSKIY, S.S.

Infrared spectra of some organotitanium compounds. Zhur. prikl.
scekt. 3 no. 61555-559 D *65 (MIRA 19:1)

1. Submitted October 6, 1964.

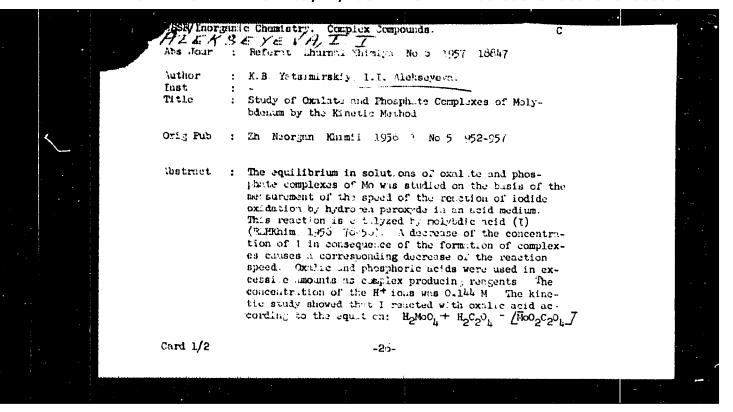


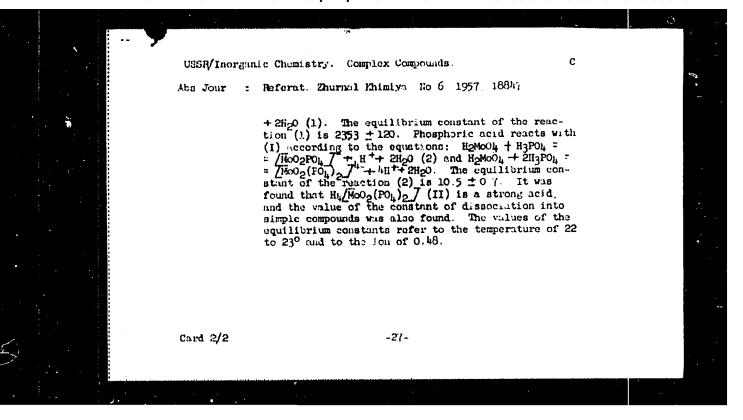


YANOVICH, T.D.; KALMYKO7A, G.N.; ALEKSEYEVA, I.K.; RACHKOVSKIY, A.P.;
CSCKINA, L.A.

Study on tuberculosis infection by means of graduated epicutaneous tuberculin test. Sbor. nauch. trud. Rost. gos. med. inst. no.22:312 '63. (MIRA 18:7)

1. Is lafedry epidemiologii Rostovskogo gosudarstvennogo meditsinskogo instituta (2av. - prof. T.D.Yanovich).





AUTHORS: Yatsimirskiy, K.B., Alekseyeva, I.I. 153 -58-1-0/29 TITLE: The Investigation of Absorption Spectra and the Determination of the Dissociation Constants of Molybdic Acid (Isucheniye spektrov pogloshoheniya i opredelaniya konstant dissotsiatsii molibdenovoy kisloty) PERIODICAL: Izvestiya vysshikh uchebnykh zavedeniy Khimiya i khimicheskaya tekhnologiya, 1958, Nr 1, pp. 53-58 (USSR) ABSTRACT: A process of depolymerization is assumed to take place during the dilution of molybdic acid solutions, and it is believed that highly diluted solutions contain nearly only monomeric particles. The correctness of this assumption is confirmed also by the study of complex molybdenum compounds (by the application of the kinetic method). The authors were therefore interested in investigating these diluted molybdic acid solutions by the optical method. They

obtained the following results: Highly diluted molybdic acid solutions contain no polymerized particles, which proves the constancy of the molar extinguishing coefficient within a fairly extensive interval of the molybdenum concentrates (2·10⁻⁵ to 2·10⁻⁴ M). By

the kinetic method it was established that in acid solutions and

Card 1/2

The Investigation of Absorption Spectra and the Determination of the Dissociation Constants of Nolybdic Acid 153-58-1-8/29

in the case of a high degree of dilution mainly non-dissociated molecules of molybdic acid exist. It may therefore be expected that in the case of low pH-values non-dissociated melecules, but at higher pH-values hydromolybdate—and molybdate fons will be found (as dissociation products of molybdic acid). For the determination of the H2McO_H-dissociation constants and for the purpose of investigating the absorption spectra the dependence of the optical density of the solutions upon the concentration of the hydrogen ions was studied. According to the course taken by the curve (fig.1) the existence of 3 kinds of particles with diverse molar extinguishing coefficients according to pH values must be expected. On the strength of the experimental data obtained absorption spectra of molybdic acid as well as of dissociation products were obtained (fig.3). There are 3 figures, 3 tables, and 6 references. 2 of which are Soviet.

ASSOCIATION:

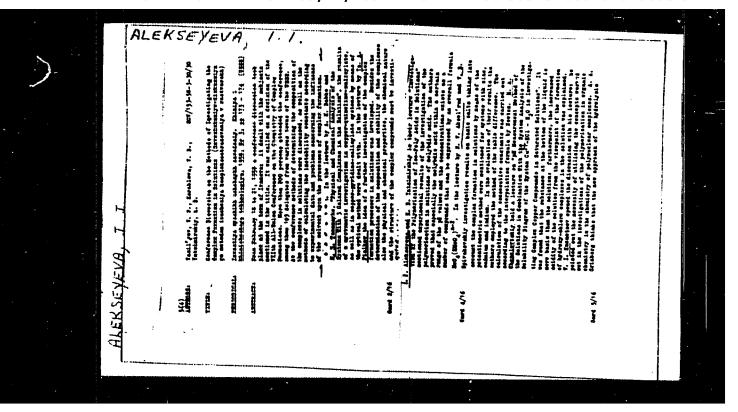
Kmfrdra analiticheskoy khimii (Chair of Inalytical

Chesistry)

SUBLITTED:

depte bor 10, 1947

Card 2/2



5(2) AUTHORS:

dataimirakiy, A.B., Alekseyeva, I.I.

SCV/32-24-12-2/45

TITLE:

Spectrophotometric and Kinetic Methods for Determining Molybdenum in Alloys (Spektrofotometricheskiy i kineticheskiy metody opre-

deleniya molibdena v splavakh)

PERIODICAL: Zavodskaya Laboratoriya, 1958, Vol 24, Nr 12, pp 1427-1429 (USSR)

ABSTRACT:

A method for determining molybdenum in steels and nickel alloys was developed which is based upon the absorption of light by molybdic acid in the wave-length region 220 - 240 mp (Ref 2). It is carried but in alkaline medium. The sensitivity of the method is given as

 7.10^{-7} g/ml Mo. The measurements were made with a SP-4 spectrophotometer. The molybdenum content is determined on the basis of a standard curve. Molybdenum alloys of Ni base Nr 3 and 4 as well as E1401 and Kh10S2M steels were analyzed (Table 1). The relative error of the methal is 2-5 %. The kinetic method of determining molybdenum is ten times mone sensitive than the spectrophotometric method. The principle of the manetic method is the catalytic effect of molybdic sold in the oxidation of indide with hydrogen percuide. The method was prevaluably reported in an earlier paper (Ref 5).

Card 1/2

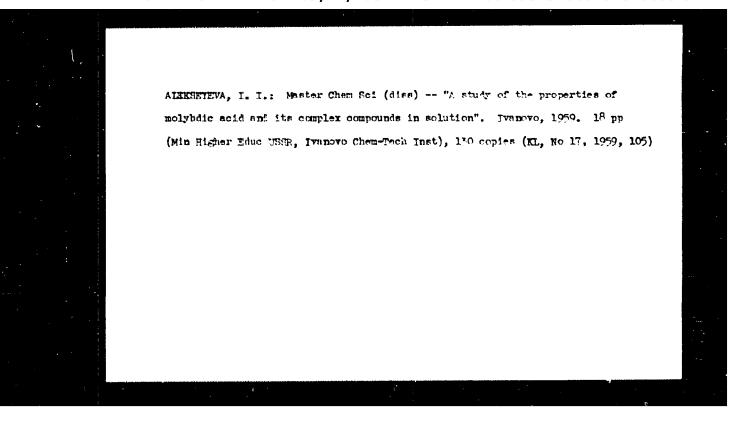
Since Fe⁵, Cu²⁺, and other ions interfere in the analysis the iron

Spectrophitemetric and Kinetic Methods for Determining Kolyidanan in Alloys

(Fa²⁺) in the present analysts was exidized with Tollin B to Fa²⁺
and then bound in a complex compound. A FKR-N appetent was used to surlyme Kh 105.2 M steel (Table 2) and iron exide (Fe₂O₃) (Table 2).

The sensitivity of the method is about 10⁻² g/ml Mo.- There are 3 tables and 7 references, 6 of which are Soviet.

ASSOCIATION: Ivanovsky khimiko-tekhnologicheskiy institut (Ivanov) Chemical Technological Institute)



5(4) AUTHORS: SUT/10

SOV/78-4-4-18/44

D

Yatsimirskiy, K. B., Alekseyeva, I. I.

TITLE:

State ? Molybdic Acid in West Acid Solutions (Solutions acidedences kisloty v solubokislykh rastvorakh)

PERIODICAL:

Zhurnal neorganicheskoy khimii, 1959, Vol 4, Nr 4, pp 8:8-822

(USSR)

ABSTRACT:

The conditions under which molybdic acid polymerizes were investigated at different concentrations (2.0.10⁻⁴ to 8.0.10⁻² molar) and at various pH values (1-6.5). In order to ascertain the possibility of the existence of the monomeric form of molybdic acid and to determine the nature of the polymerization process the light absorption of solutions of molybdates and molybdic acid was studied at wavelengths from 270 to 350 mm. The determination of the optical densities was carried out using a SF-4 spectrophotometer. The dependence of the molar absorption coefficients upon the molybdate concentration was investigated at constant pH and constant wavelength. The curvacture is show that at lower molybdate concentrations the molybdic acid is present in the monomeric form. The polymeric form exists also at higher concentrations. The phase diagram of molybdic acid in solution was constructed (log C in

Card 1/2

SOV/78-4-4-18/44

State Nolybdic Acid in Weakly Apid Solution

dependence upon the pH value). In the pH range i to 4 the polymerization begins at a molybdic acid concentration of 10 molar. With an increase in the pH value the polymerization boundary shifts to the side of the higher concentration. Finally at pH 6.5 no more polymerization occurs. The following polymerization schemes are suggested for molybdic acid:

 $nMoO_4^{2-} + (n-1)HOH = (HMoO_4)_{n-1}MoO_4^{(n+1)-} + (n-1)OH^-$

(at pH values 4 to 6) and

 $(n+1)H^{T} + (HMoO_{4})_{n-1}MoO_{4}^{(n+1)} \rightleftharpoons (H_{2}MoO_{4})_{r}$

(at pH values 1-4).

There are 5 figures and 7 references, 1 of which is Soviet.

ASSOCIATION:

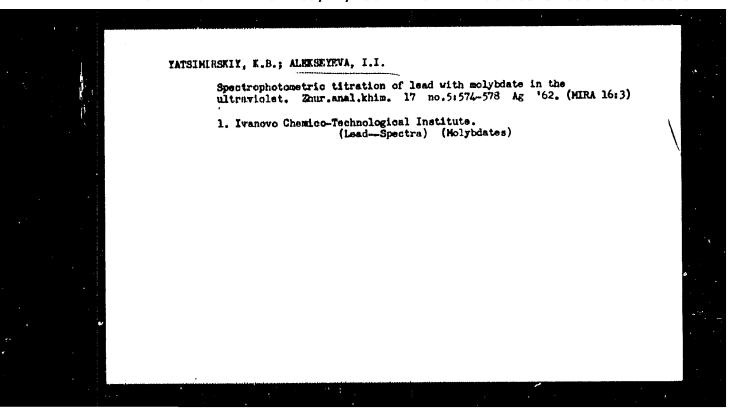
Ivanovskiy khimiko-tekhnologicheskiy institut (Ivanovo Chemi; x2

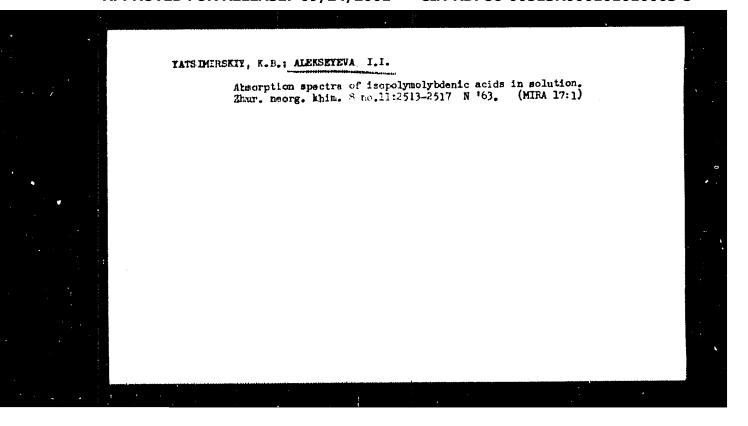
Technological Institute)

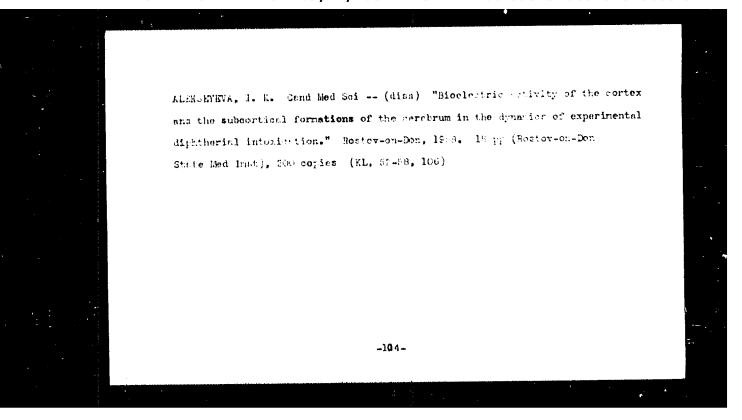
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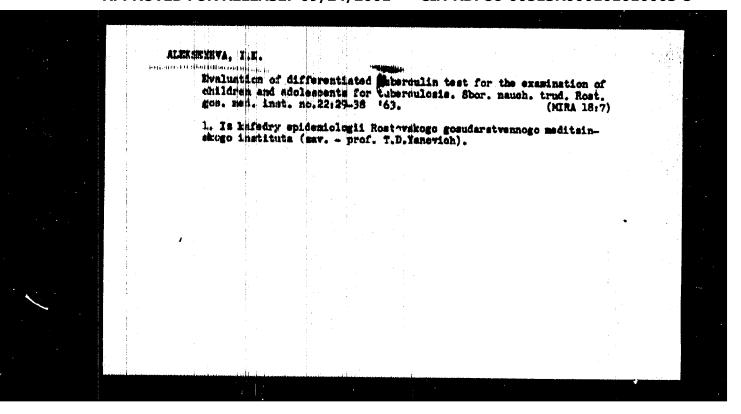
April 24, 1958

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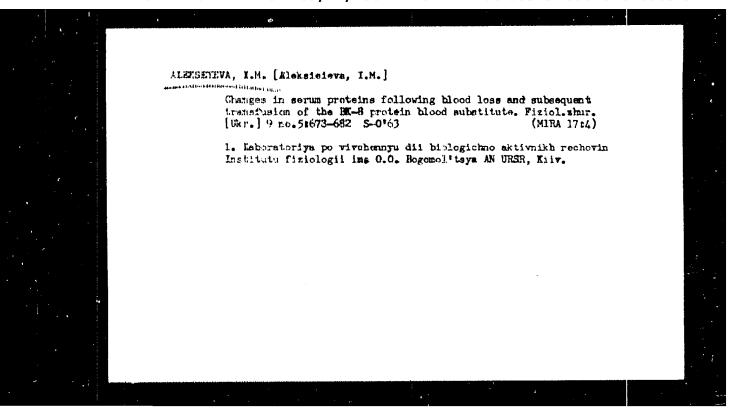
ALEKSEYEVA, I.M. [Aleksieieva, I.M.]

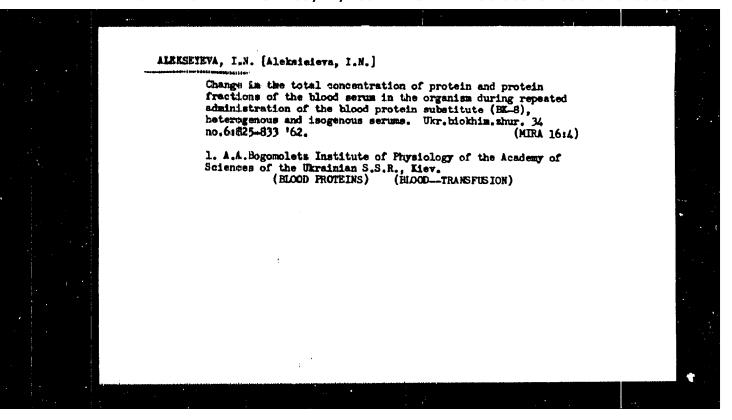
Correlation between protein changes in the blood serum and variation of the arterial pressure in heterohemotransfusion shock. Fiziol. zhur. [Ukr.] 11 no.1:121-123 Jn-F '65. (MIRA 18:7)

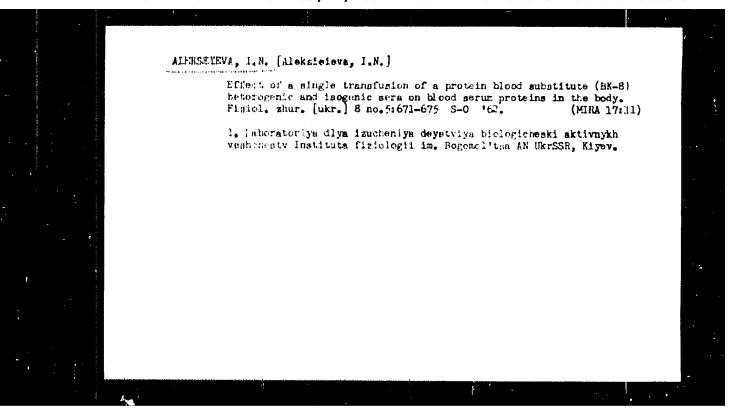
l. Laboratoriya izuchaniya biologichaski avtivnykh veshchestv Instituta fiziologii im. Gogomolitsa AN UkrSSR, Kiyev.

L 31282-66 ACC NR. AP6020241 SOURCE CODE: UR/0300/65/037/002/0226/0230 AUTHOR: Aleksynyeva I. M. 19 ORG: Institute of Physiology im. A. A. Bogomolets, AN UkrSSR, Kiev (Instytut fiziolohiyi AN UkrSSR) Ŀ TITLE: Protein changes in the serum of the recipient after transfusion of a protein blood substitute (BK-8) 22 SOURCE: Ukrayins kyy biokhimichnyy zhurnal, v. 37, no. 2, 1965, 226-230 TOPIC TAGS: protein, blood, serum ABSTRACT: Previous investigations of the author have shown that transfusion of BK-8 [belkovyy krovosamenitel' -8; protein blood substitute-8] induces a decrease in the albumin content and an increase in that of alpha- and beta-globuling during the first five minutes - one hour after the transfusion. This report deals with studies determining protein composition of blood serum five minutes and one hour after transfusion of BK-8 under conditions of precipitation of BK-8 proteins changes noted in the protein formula disappear. Therefore, it can be assumed that the changes in the protein content of the serum - the decrease in albumin content and increase in alpha- and beta-globulin content - are associated in the early post-transfusion period with the transformation of protein fractions due to the transfer by them of some of the blood substitute proteins. Orig. art. has: 2 tables. [JPRS] SUB CODE: 06 / SUBM DATE: 07Jan64 / ORIG REF: 005 / OTH REF: 001 Card 1/1 / //

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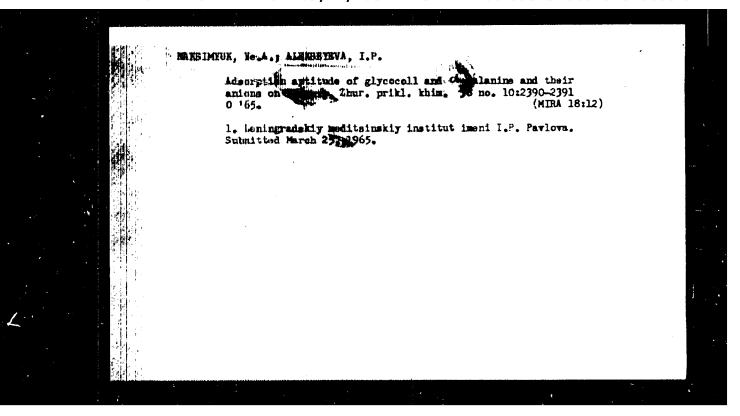




SPASORURGHENT, Yu. A. [Spasokukote kyi, Ru. 0.]; ALEMSEYEVA, I. N. [Aleksiefeva, I. N.]; FERCHOVERATA, N.I. [Perconvertan, R.T.]

Effect of intravenous injections of high comes of antievarial cytotoxic serum on the secural cycle of whate rats, Pizicl. whur. [Ukr] 9 no. 3: 393-324 htmlere '63. (KIRA 18:1)

1. Laboratoriya Izueneniya biologicheski uktivuyhh vesmehestv Instituta fiziclogii iz. Begonol'tas AN UkrSCR, Kiyev.



s/079/60/030/007/021/039/xx B001/B066

AUTHORS:

Alekseyeva, I. P., Pesotskaya, V. M., and Ptitsyn, B. V.

TITLE

Oxidation Potential of Permanganate

PERIODICAL: Zhurnal obshchey khimii, 1960, Vol. 30, No. 7, pp. 2104-2108

TRIT: The purpose of this work was a more exact characterization of the oxidizing effect of permanganate. Its oxidizing effect depends on the acidity of the solution, but research workers disagree in this respect. The investigation of the dependence of the potential of the permanganate solution on pH is of interest since the reduced form which is in equilibrium with the permanganate ion may be determined from the potential as a function of the percentual content. In an acid solution, the latter may be in equilibrium with some of its reduction products, except compounds of bivalent manganese, as potassium permanganate oxidizes the Mn. at any percentual content. Therefore, the universal equation Mn0'4 R° 3

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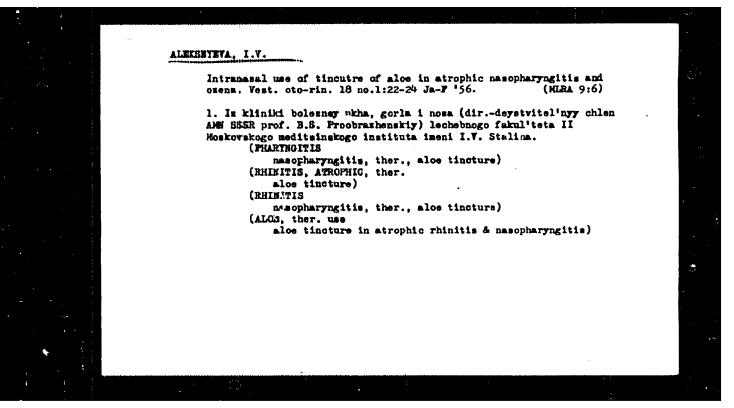
Oxidation Potential of Permanganate

S/079/60/030/007/021/039/XX B001/B066

cannot be used to characterize the oridation properties of permanganate. To study these states of equilibrium different methods were applied (Refs. 3-5,6.7). B. V. Ptitsyn and V. F. Petrov (Ref. 8) indicated that also in the case of bichromate solution the oxidation potential may be affected by the nature of the acid. This might also apply to permanganate. The present paper deals with the following problems: 1) Quantitative determination of the oxidation potential of the permanganate solution as dependent on the pH of the medium. 2) The question as to whether the nature of the acid influences the potential. 3) A more exact characterization of the states of equilibrium established in the permanganate solution under the conditions of the experiments to be performed. The authors investigated the defendence of a potassium permanganate solution (0.01 mole) in solutions of HClO₄, HNO₃, H₂SO₄, CH₃COOH. The effect depends on the nature of the acid at low pH only. An attempt was made to interpret the nature of the reduced forms which are in equilibrium with the permanganate ion, on the basis of the functions of the permanganate solution potential on pH in solutions of different acids. Table 1 gives the data of the potential dependence of KMnO₄ dissolved in HClO₃, and Table 2 the values

Card 2/5

Ourdation	Potential of Permanganate	\$/079/60/030/007/021/039/23 B001/B066				
		agram illustrates the oxidation expendent on pH in the presence of tial of permanganate in HClO, and				
HNO _x , E va	s found to be 1600 mv, in Haso	1 E = 1650 my. There are				
HNO_3 , E was found to be 1600 mv, in H_2SO_4 : E = 1650 mv. There are 1 figure, 2 tables, and 9 references: 4 Soviet, 4 US, and 1 Spanish						
	ASSOCIATION: Leningradskiy tekhnologicheskiy institut pishchevoy promyshlennosti (Leningrad Technological Institute of the Alimentary Industry)					
SUBMITTED:	July 6, 1959					
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Card 3/3						
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REZNIKOV, L.I.; SHAMASH, S.Ya.; ALEKSEYEVA, I.V.

State of students' knowledge in physics. Fig.v shkole 21
no.4150-53 Jl-Ag '61. (MIRA 14:10)

1. Sektor obucheniya fiziko Instituta obshahego i politekhmicheskogo obrasovaniya Akademii pedagogicheskikh nmuk RSFSR.

(Physics—Study and teaching)

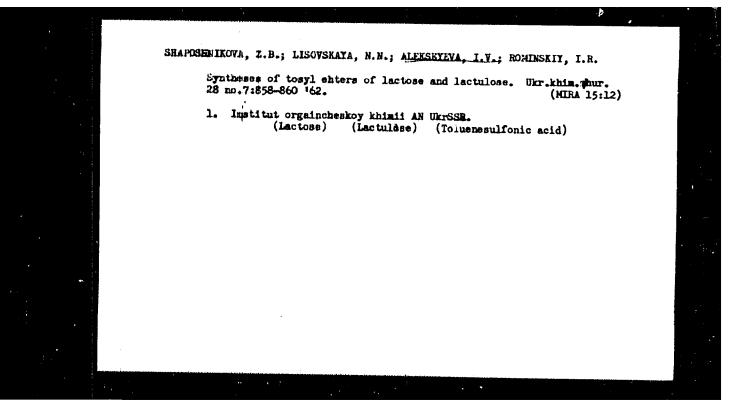
SHAPOSHEIKOVA, 2. B.; ALRESETEVA, I. V.; ROMINSKIY, I. R.

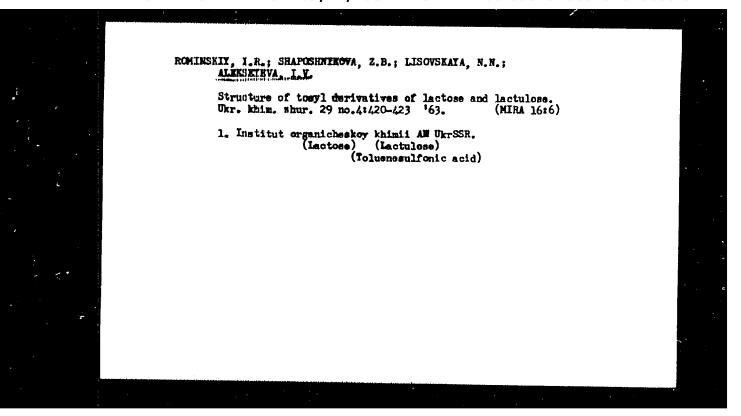
Mathod of preparing pure lactulose with the aid of ion exchange resins. Ukr. khim. shur. 28 no.6:724-725

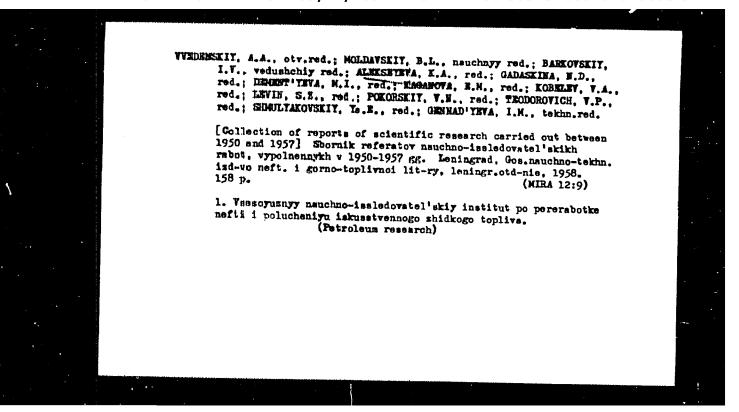
(MIRA 15:10)

1. Institut organicheskov khimii AN UkrSSR.

(Lactulose) (Ion exchange resins)







SCT/65-58-6-4/13

AUTHORS: TITLE:

Rudkiwskir, D. M; Trifel', A. G. and Alekseyera, K. A.

Preparation of $C_6 = C_8$ Alcohols from Olefin-Jontaining Puel Fractions by the Oxo-Synthesis. (Polucheniye Tysshikh spirtor $C_6 = C_8$ is elefins: derehash shikh toplitnykh fraktsiy setedat eks.sietera).

PERIODICAL:

Khimiya i Tokumulikiya Toplis i Masel. 1959. Nr.6. pp. 17 . 24. (OSSR).

ABSTRACT:

The fundamental principles of the Org-synthesis and the uses of and products one recieved. Arongst these end products are higher alreads (C_6,C_8) which are excellent fictation agents for light metal tree and for slack. Results of experiments in the preparation of C_6-C_8 aldebrases. hydes from elefin-centaining fuel fractions are given. The influence of the concentration of the satalyst, the temperature, pressure, composition of the synthesis gas, the rate of supply of the liquid raw material and of the rate of directables of the gas on the carbonylation process, were investigated. The raw material used was the fraction boiling up to 100 7 which was separated on a restification sclumn during two distillations of bracked petroleum. Various physical constants of this fraction are listed as well as the content of 05. 06 and 07 hydro-

Card 1/4

50% 65-58-6-4/13

P reparation of Co + Cg Alcohols from Olsfin-Containing Fuel Fractions by the Oxc-Synthesis

were carried out on a continuous apparatus (Fig.1). Notails of the process of carbonylation of unsaturated hydrocarbons are given. Cobalt carbonyl was used as catalyst. The influence of the concentration of this catalyst on the rate of carbonylation of unsaturated Cg - Cy hydrocarbons was investigated at a temperature of 16200, pressures of 200 and 300 atms and the ratic of the rate of supply to the raw material was 3.6:1. The volume of circulating gas = 0.7m / itms of raw material. The concentration of the catalyst was changed within the limits 0.03 - 0.315. Results of these experiments are given in Table 2 and Fig.2. Details of investigations on the influence of temperature on the rate of the reaction at 290 atms are given in Table 3 and Fig.3. Activation energy was calculated according to the equation by Arrhenius and was - 11.000 cals mole. The temperature coefficient of the rate of reaction = 1.4. Experiments on the effect of pressure on the carbonylation process were carried out at liw depths of conversion (Table 4 and Fig.4.). When the reaction was carried out under

Card 2/4

SCM/65-58-6-4/13 Preparation of $C_6 = C_6$ Alsohels from Olefin-Centaining Puel Practions by the Oxo-Synthesis.

industrial conditions (volume rate # 3, and concentration of the satalyst * 0.8%) a change in the pressure from 150 - 300 atms does not affect the depths of concersion (Table 5). Indescigations on the influence of the composition of the gas on the process were partial out at varying temperatures, partial pressures of 60 and H2 and varying rates of supply of the raw material. From data given in Table (and Pig.5 it can be seen that at low temperatures (120 - 140 0) the depth of conversion of unsaturated lydrocarbons increases with increasing partial pressure of hydr ger. Hasults of tests on the influence of the mate of supply of the raw material and the quantity of dr ulating gas on the carbonylation process are given in Tables 7 and 6. The analytical imposingations showed that the products obtained from fractions up to 10000 contain 39%-48% exygen-containing impounds. The alcohols were reparated from the hydrogenates by rectification; the fraction booking up to 10000 (unreasted raw material); the alcohol fraction to the notion of the constants of all these fractions are given. There are 5 Tables, 5 Figures,

Card 3/4

SOW/85-58-6-4/13

Preparation of C6 - C8 Alochels from Olefin-Containing Fuel Practions by the Oxo-Synthesis.

and 5 References: 3 Sowiet, 1 German and 1 English.

ASSOCIATION: Lennii

Card 4/4

AUTHORS:

SOV/65-58-10-2/15

Alekseyeva, K. A. and Moldavskiy, B.

TITLE:

Conversion of Oxygen-Containing Compounds of Charachave Tar During Liquid-Phase Hydrogenation (Prevrashcheniye kislorodsoderzhashchikh soyedineniy Cheremkhovskoy

smoly pri zhidkofasnoy gidrogenizatsii)

PERIODICAL: Khimiya i Tekhnologiya Topliv i Masel, 1958, Nr 10, pp 7 - 11 (USSR)

ABSTRACT:

Results of investigations on the conversion of orygen-containing compounds are reported as most publications on this subject deal with the selection of catalysts and conditions of reduction of the oxygen-containing compounds, especially of phenols into hydrocarbons (Ref.1). Tar from Cherenthovo 's semi-coke was tested. The degree of conversion was evaluated according to the formation of the broad fraction (the fraction of the hydrogenation product boiling up to a temperature of 325°C). This fraction was freed of its light components by vacuum distillation. Physical data are tabulated. The experiments were carried out in an apparatus (Fig.1) which comprised a 2 litre isothermic autoclave with an electromagnetic drive and an agitator which was operated

Card 1/4

SOV/65-58-10-2/15 Conversion of Oxygen-Containing Compounds of Characteristic Tar Puring Liquid-Phase Hydrogenation

at 2,800 rers/minute. S. A. Babushkin's method (Ref.5) was used with slight modifications for the direct determination of oxygen in the organic substances. The oxygen content in the initial tar and in the hydrogenation products (the broad fraction, the residue above 325°C, the gas and water) defined also the degree of conversion of the oxygen-centaining substances. The influence of the length of the reaction, the temperature and pressure on the destructive hydrogenation of Cherenthare tar was also investigated. A series of experiments was carried out at a pressure of 290 atms and temperatures of 460 to 420°C (Table 1 and Fig.2). The highest degree of decomposition of the raw material occurs at the beginning of the reaction, after 20 minutes, when the temperature equals 460°C, and after 1 hour at a temperature of 420°C. The rate of conversion of oxygen-containing compounds is much quicker. The effect of the pressure of hydrogen on the conversion of oxygen-containing compounds was tested at a temperature of 460°C when the reaction was carried out for one hour at pressures of 290, 200 and 100 atms. A change in the pressure from 290 to 100 atms hardly affected the total

Card 2/4

Conversion of Oxygen-Containing Compounds of Cherenthore Tar During Liquid-Phase Hydrogenation

conversion of the oxygen-centaining compounds which varied within the limits of 84 to 85%, but led to the distribution of the exygen between the broad fraction, water and gas. Hydrogenation of Chereskhovo tar at 100 atms pressure is accompanied by coke formation, and the residue beiling above 325°C shows low hydrogen content and considerable increase in the specific weight of the residue. This raw material should not be hydrogen mated at 100 atms pressure in the presence of low-activity catalysts such as iron. The temperature coefficients for various degrees of conversions were also calculated (Table 3). Under the given process conditions, the decomposition of high molecular oxygen-containing compounds results in the formation of lower phenols and neutral oxygen-containing compounds having boiling temperatures in the range of those of the broad fraction: Table 4. An increase in the temperature intensifies the decomposition process of the raw material and also the conversion of the oxygen-containing compounds. The amount of phenols and neutral oxygen-centaining compounds increases only slightly on raising the temperature because

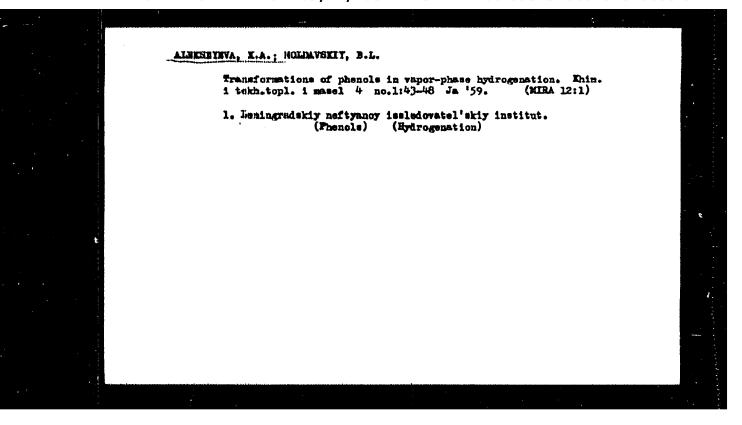
Card 3/4

Conversion of Oxygen-Containing Compounds of Cherenkhove Tar During Liquid-Phase Hydrogenation

under the given condition the rate of formation of compounds only slightly exceeds the rate of their reduction. There are 4 Tables, 2 Figures and 7 Soviet References.

ASSOCIATION: LenNII

Card 4/4



AUTHORS: Rudkqvskiy, D. M., Trifel', A. G., S/064/59/000/08/02/021 B115/B017

TITLE: Production of <u>Butyric Aldehydes</u> and <u>Butyl Alcohols</u> by Means of the Method of Oxosynthesis

PERIODICAL: Khimicheskaya promyshlennost', 1959, Nr 8, pp 652-658 (USSR)

ABSTRACT:

In the present paper the production of butyric aldehydes and butyl alcohols from a commercial propane - propylene fraction and from a carbon monoxide - hydrogen mixture by means of oxosynthesis is described, and the technological factors determining this process are investigated. The method has been described already earlier (Ref 7). It consists of three stages: production of the cobalt-carbonyl solution (which is used as catalyst, solvent: toluene, iso- and n-butyl alcohol, pentane-hexane fraction from the direct distillation of gasoline), carbonylisation and decomposition of the catalytic complex formed. The apparatus used and the processes which take place in them are briefly described. Figure 1 shows the scheme of the laboratory arrangement, in which a flow system was used and work was carried out at a temperature of approximately

used and work was carried out at a temperature of approximately 150° and at pressures of 150 to 300 atm. The composition of the Card 1/3 gases used as initial products is also given. The influence exer-

Production of Butyric Aldehydes and Butyl Alcohols by Means of the Hethod of Oxosynthesis

Card 2/3

S/064/59/000/08/02/021 B115/B017

cised by the temperatures in the range of from 110 to 150° on the rate of carbonylisation of propylene is investigated in a static system. The following was also investigated: The influence exercised by the cobalt concentration on the conversion of propylene at 120, 135 and 150° and 150 atm (Fig 3), the influence of pressure on the carbonylisation of propylene (Table 1), of the propylene concentration in the solution on the carbonylisation of propylene (Table 2), of the gas composition on the rate of pentane carbonylisation (Fig 4), of propylene (Table 3) at different temperatures, of the ratio P_{CO}: P_R, on the constant of reaction rate (K-10²) (Fig 5), of the partial pressure of carbon monoxide P_{CO} on the maximum stability temperature of cobalt carbonyl (Fig 6), of the composition of the propane-propylene fraction (Fig 7) and of the volume rate of the liquid raw material (Table 4) on the yield in propylene transformation products. Carbon dioxide delays the carbonylisation reaction. The maximum stability temperature of cobalt carbonyl shows a logarithmic dependence on the partial pressure of carbon monoxide. The influence exerted by various factors on the formation of acetals in the condensation products in using butyl alcohols as solvent is given (Table 5), and the

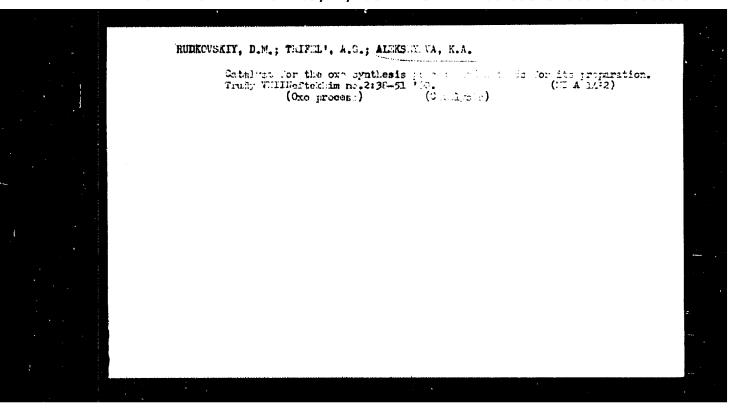
Production of Butyric Aldehydes and Butyl Alcohols

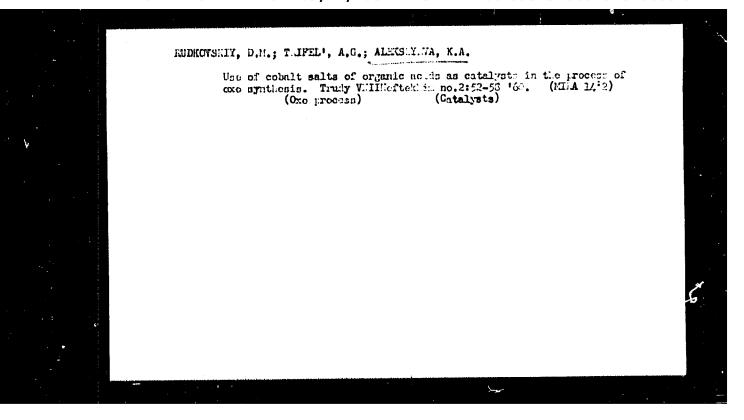
by Means of the Method of Oxosynthesis

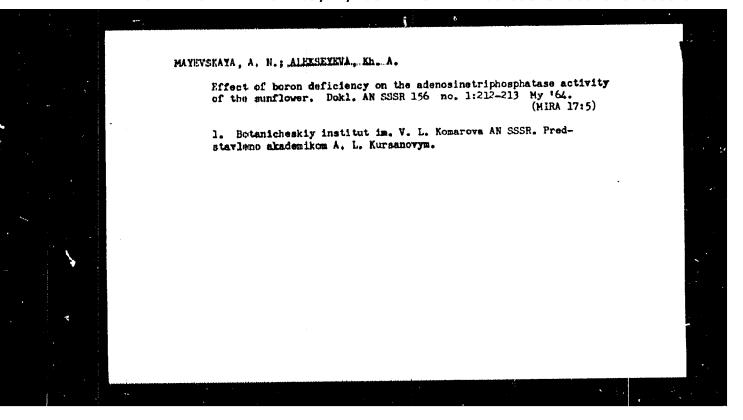
Composition of the hydrogenated product obtained by using a
pentame-hexane fraction as solvent in the carbonylisation of the
propame-propylene fraction is mentioned (Fig 8, Table 6). The results show that n-butyl alcohol is the main reaction product
(60%). The other products are: isobutyl alcohol (22%), alcohols
C8 and ester (6%), 2-ethyl hexanol (9.5%), and higher condensation
products (higher than C8) (4%). There are 8 figures, 6 tables, and
11 references, 4 of which are Soviet.

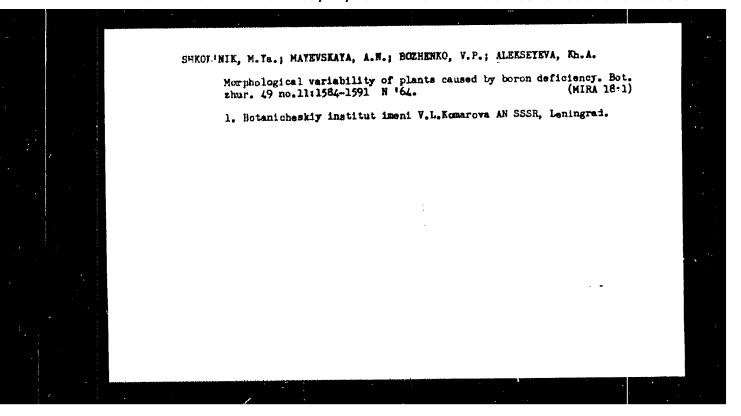
ASSOCIATION: VHIIneftekhim (VHIIneftekhim - All-Union Scientific Research
Institute of Petroleum Chemistry)

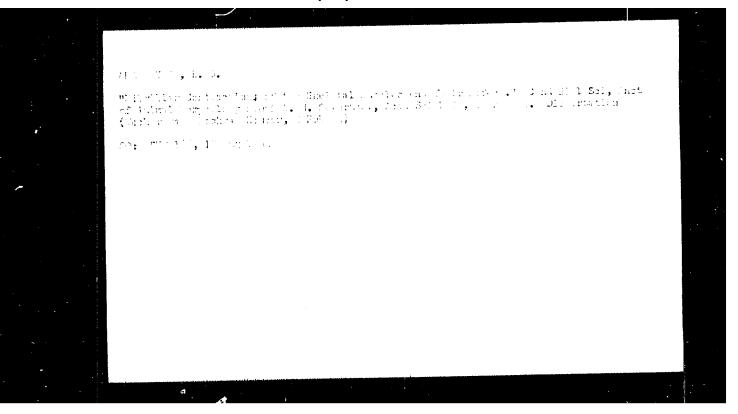
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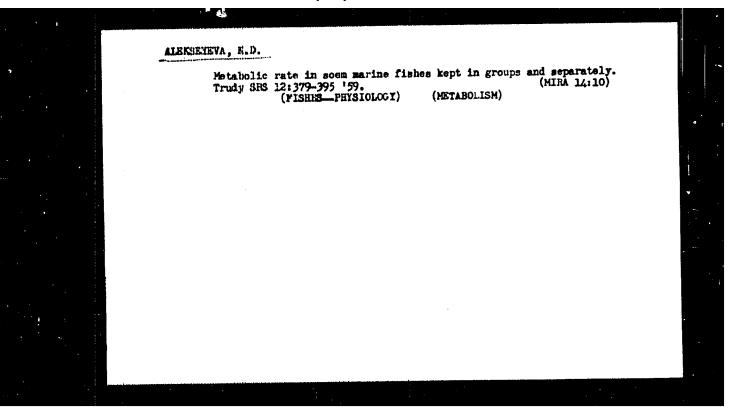




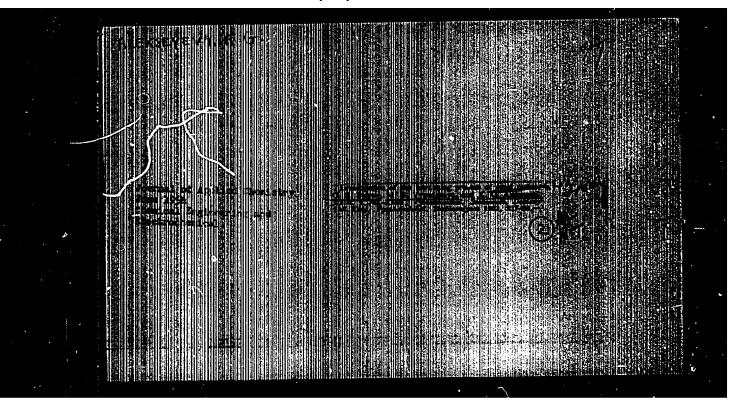






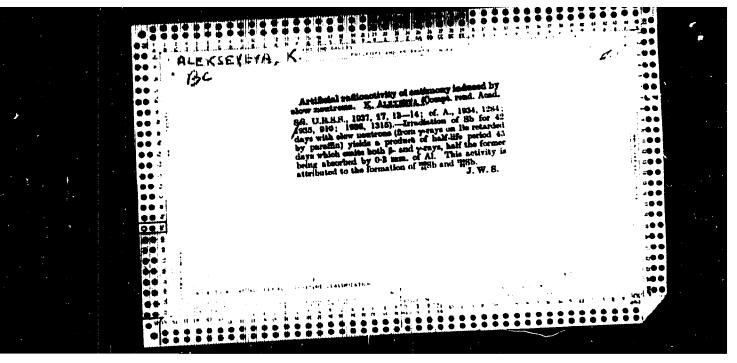


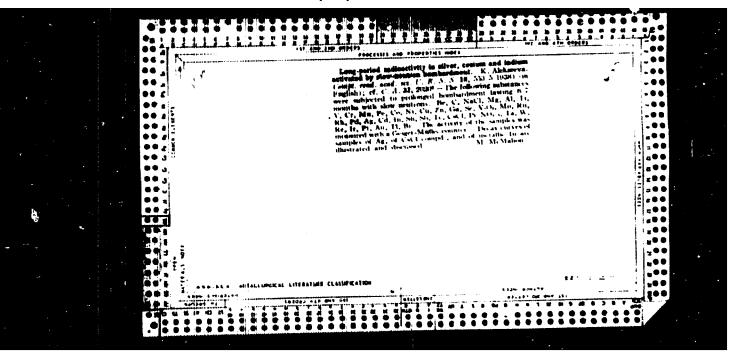
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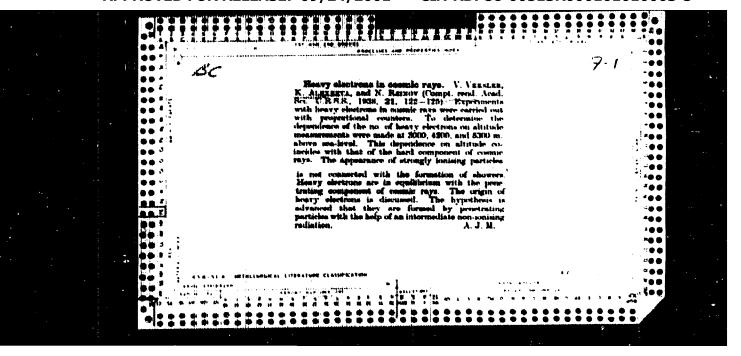


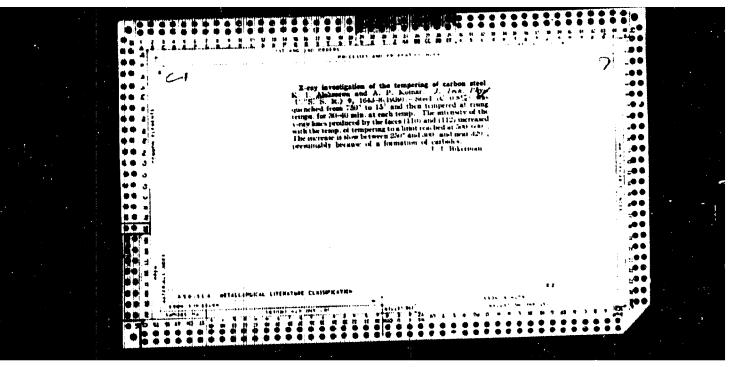
*Roentgenography and its Application to the Study of Materials."

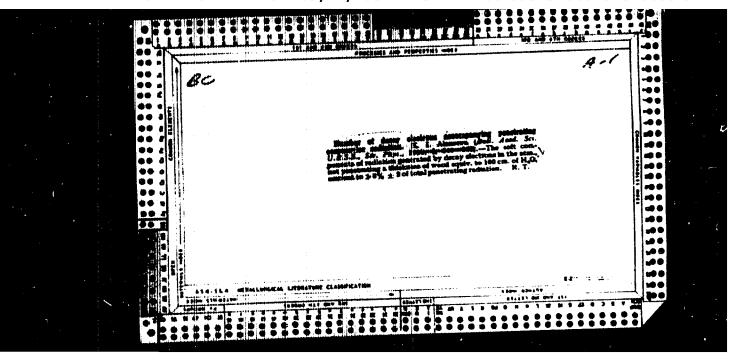
ZhTF 5, 342, 1935

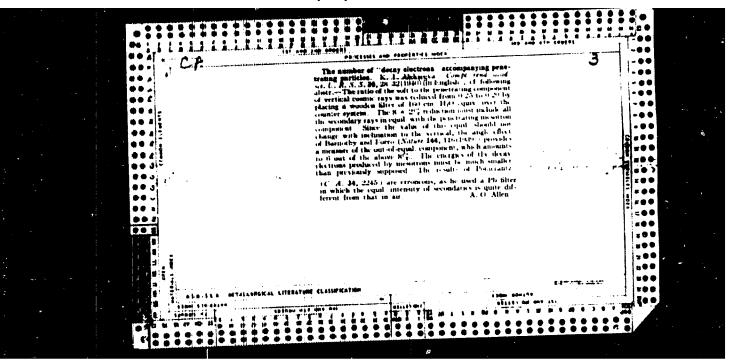


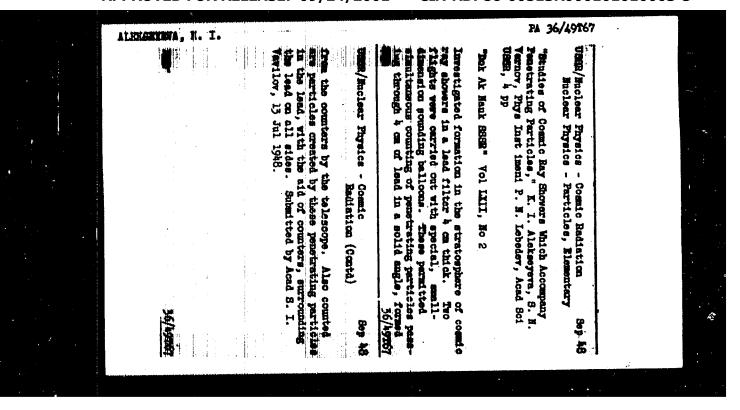


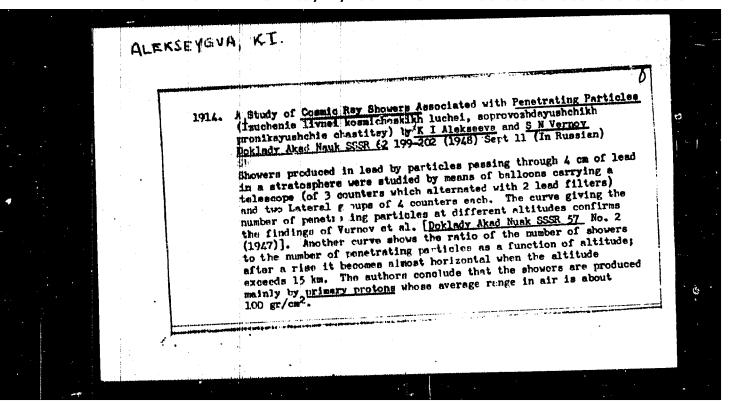


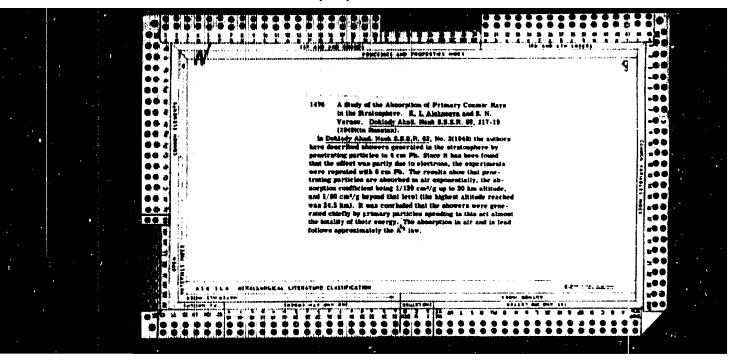


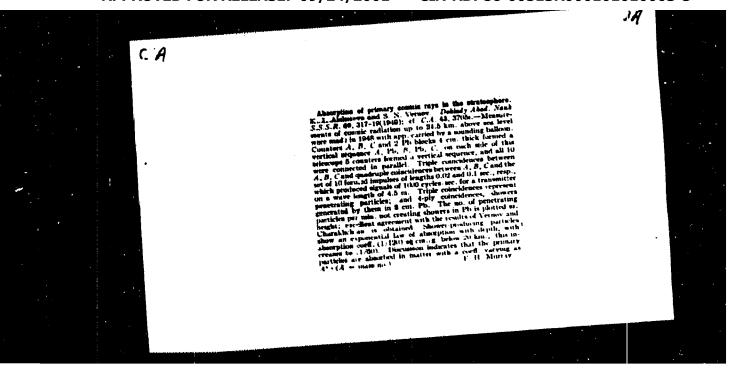


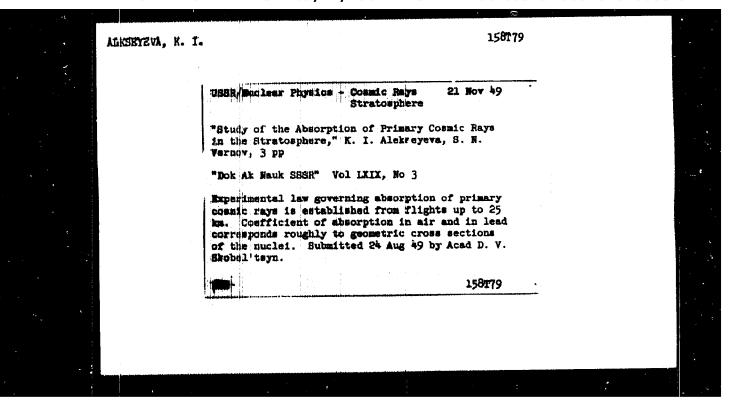


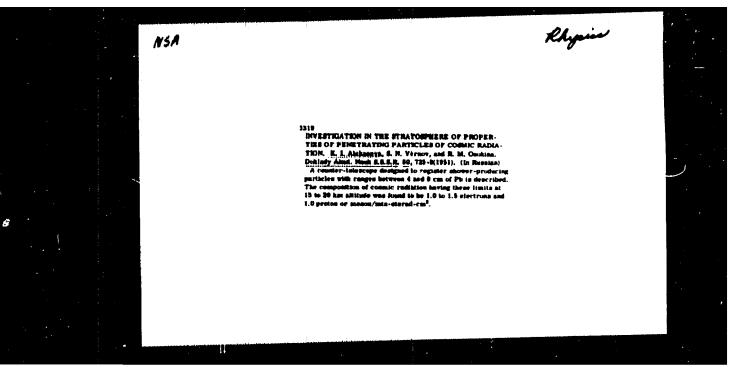






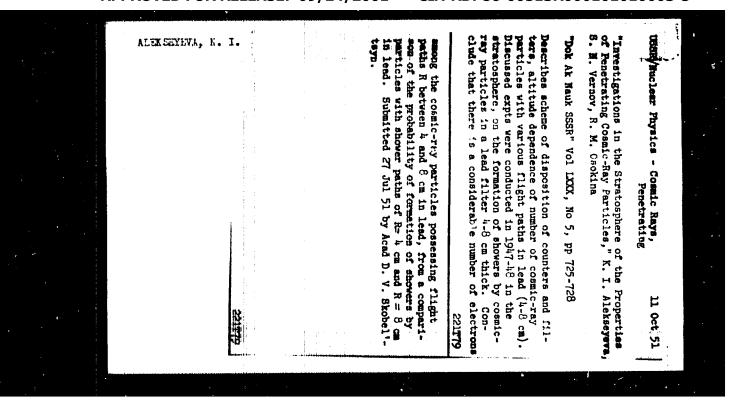






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CIA-RDP86-00513R000101010003-5



AUTHOR TITLE

ALEXNETEVA, K.I., GRIGGROV, N.L.,

PA - 2713
The Inelastic Interaction of Protons with an Energy of more than 7 BeV with Carbon- and Hydrogen-Nuclei.

(Neuprugoye vzaimodeystviye protonov s emergiyey bol'she 7 BeV s

yadrami ugleroda i vodoroda - Russian)

Zhurnal Eksperim. i Teoret. Fiziki, 1957, Vol 32, Nr 1, PERIODICAL

pp holi-ho5 (U.S.S.R.) Received 5/1957

Reviewed 6/1957

ABSTRACT

Card 1/2

The gross section of this inelastic interaction was measured in the stratosphere in an altitude of from 20 to 25 km in the geomagnetic latitude 31, where the minimum kinetic energy of the primary protons amounts to 7 NeV and the energy of 60 of all protons amounts to from 7 to 20 BeV. The measuring arrangement, i.e. the distribution of the telescope, filter, and counter is discussed. The cross section of the inelastic protons with carbon nuclei was determined by two methods. 1) From the reduction of the flow of the individual shower-forming particles which, in the case of the existence of graphite in the telescope, drop on to the lead filter. This reduction is due to inelastic acts of interaction of the protons with the atomic nuclei of the carbon. (Measurements were taken of the modification of the number of electron-mucleus showers coming out of the lead in the case of the existence of graphite in the telescope).

2) By direct measuring of the number of the electron-nucleus showers

produced in the graphite.

The Inelastic Interaction of Protons with an Energy of more PA-2713 than 7 BeV with Carbon- and Hydrogen-Nuclei.

By method 1) the authors obtained the following values of the range and theoress section of the inelastic interaction of the protons with carbon nuclei $L_p^C = 67_{-9}^{+13} \, g/cm^a$, $\sigma_p^C = 300 \pm 50$ millibarn.

The necessary corrections for these values have already been taken into account. By method 2) they obtained values $_{\rm LP}^{\rm C}$ =7.3 + 7 g/cm² of = 270 + 30 millibarn. The error given for these values in all cases denotes the statistical error. The authors determined the cross section of the inelastic proton-proton interaction from the difference of the mumber of the electron-nucleon showers recorded in paraffin and graphite and they hereby obtained the following values $_{\rm LP}^{\rm H} = 1.7^{+37}$ g/cm² of = 35 + 16 millibarn. This value of the cross sectionwill—probably not change by more than lo/o, if the 5 shower is taken into account. At present the authors are investigating the production of 5 showers at sea level in paraffin and graphite. (No ill.)

ASSOCIATION PRESENTED BY SUBMITTED AVAILABLE

Card 2/2

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ALEKSTYEUM ベエ 20-1-18/54 Alekseyeva, K.I., Brikker, S.I., Grigorov, N.L, Savin, P.D., AUTHOR: Shcherbakov, N.A. Determination of the Flux of Primary Cosmic Ray Particles at TITLE: 31 N. Latitude (Cpredeleniye potoka perwichnykh kosmicheskikh chastits na shirote 510 N.) Doklady Akademii Nauk SSSR, 1957, Vol. 115, Nr 1, pp. 71 - 74 PERIODICAL: (USSR) On the occasion of the determination of the cross-section of ABSTRACT: the non-elastic interaction of the cosmic ray particles with the nuclei of carbon and hydrogen the authors determined the intensity of the hard components in stratosphere. These measurements made possible the determination of the flux of the cosmic particles at the border of atmosphere. The authors describe the apparatus for the measurements of the cross-section of nonelastic interaction as well as the results of the measurements of the intensity of hard components. (The results of the measurements of the cross-sections shall be published in later works). The scheme for the arrangement of the filters as well as of the counter is shown by means of a sketch. The telescope and the Card 1/3 filters were surrounded by many hodoscopic counters for the re-

20-1-18/54

Determine tion of the Flux of Primary Cosmic Ray Particles at 31 °K. Latitude

gistration of secondary particles. Furthermore three series of telescopic counters were connected in form of a hodoscope. The inpulse for the control of this apparatus was the triple coincidence in the three series of counter. All results obtained with these apparatuses were transmitted to the ground by radio. The elaboration of the measurement results obtained this way are discussed for the following cases: single particle, shower which developed outside the filter, shower in the upper part of the apparatus and shower in the lower part of the apparatus. The course of particles as a function of altitude which cause no interactions in a Pb and Al filter is shown by a diagram. Another curve shows the number of the nuclear interactions in Pb and Al filter. Another curve is the sum of the two mentioned curves, that is to say, it characterizes the total flux of the particles of the hard component at various altitudes. This flux is 2,0 particles/cm mm. sterad at the border of atmosphere. In the end the results obtained are compared with those of other authors. There are 2 figures.

Card 2/3

Determination of the Plux of Primary Cosmic Ray Particles at 31° N. Latitude

ASSOCIATION:

Moscow State University im. M. V. Lomonosov
(Moskovskiy gosudarstvennyy universitet im. M.V. Lomonosova)

PRESENTED BY:

Junuary 12, 1957

AVAILABLE:

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Card 3/3

ALERSE YEUR XI 20-2-17/62 AUTHORS: Alekseyeva, K.I., Grigorov, N.L. TIPLE The Cross Section of the Inelastic Interaction of More Than 7 BeV Kimetic Energy Protons with the Atomic Nuclei of Carbon (Secheniye neuprugogo vzaimodeystviya protonov s kineticheskoy energiyey bol'she 7 BeV s atomnymi yadrami ugleroda) PERIODICAL: Doklady Akademii Nauk SSSR, 1957, Vol. 115, Nr 2, pp. 259 - 262 (USSR) The present paper gives the data of the measurement of the cross ABSTRACT: section of the inelastic interaction of protons with carbon nuclei on the geomagnetic latitude 31° in the stratosphere in the domain of those energies for which the authors earlier found a small non-elasticity of the collision with light nuclei. The apparatus for the measurement of this inelastic interaction consisted of a telescope surrounded by many hodoscopic counters. 80 % of the proton current had a kinetic energy of more than 7 BeV and 3/4 of these had the average energy \sim 12 BeV. The cross section of the non-elastical interaction proton-carbon nucleus Card 1/3 was determined by 2 methods:

The Cross Section of the Inelastic Interaction of More Than 7 BeV Kinetic Energy Protons with the Atomic Muclei of Carbon 1.) By the method of reducing the flow of the shower-forming component during the passage through graphite: The method does not require the knowledge of the true intensity of the flow of the shower-forming particles and of the true number of the interactions with lead. For the range obtained by this me thod in 20-25 km altitude resp. for the interaction of the inelastic interaction of the protons and the carbon nuclei $L_{\rm p}^{\rm C} = 63_{-9}^{+12} \, \rm g \cdot cm^{-2}$ resp. $\sigma_{p}^{C} = 315 \pm 50 \text{ mb is found.}$ 2.) By the method of direct observation of the inelastic interaction in a graphite filter (from the number of showers proceeding from the mesotrons produced in graphite) the authors determined the values $L_p^C = 76 \pm 7 \text{ g.dm}^{-2}$, $\sigma_p^C = 264 \pm 24 \text{ mb}$. Suppositions on the cause of the differences between the values found by both methods are given. Conclusions: The here-measured cross section of the inelastic interaction of the protons of the mean energy \sim 12 BeV with the Card 2/3 carbon nuclei amounts to 315 \pm 50 mb, thus it agrees within the D

20-2-17/62

The Cross Section of the Inelastic Interaction of More Than 7 BeV Kinetic Energy Protons with the Atomic Nuclei of Carbon

limits of error with the geometric cross section of the carbon atom. There are 2 figures, 2 tables, 9 references, 3 of which are

Slavic.

ASSOCIATION: Moscow State University ineni M.V. Lomonosov

(Noskovskiy gosudarstvennyy universitet imeni M.V. Lomonosova)

PRESENTED:

January 15, 1957, by D.B. Skobel'tsyn, Academician

SUBMITTED:

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January 12, 1957

AVAILABLE:

Library of Congress

Card 3/3

ALEKSEYEVA. K.I. 20-4-13/52 AUTHORS: Alekseyeva, K. I., Grigorov, N. L. Proton-Proton Interaction at The Cross Section of non-Elastic TITLE: Proton Energies Exceeding 7 Bev. (Secheniye neuprugogo vzaimodeyatvi= ya proton-proton pri energiyakh protonov bol'she 7 BeV). Doklady AN SSSR, 1957, Vol. 117, Wr L, pp. 593-596 (USSR). PERIODICAL: First a brief report is given on 3 previous works dealing with this ABSTRACT: subject. The authors applied the data obtained from flights with two devices with lead filter and one apparatus without lead filter for the determination of the cross-section referred to in the title. On order to make the conditions of registration of the showers develor ping both in paraffin and graphite approximately equal, the authors used graphite in form of powder. The necessary corrections of the measurements are discussed as well. The authors computed the crosssection of inelastic proton proton interaction from the ratio of the streams of the individual shower-developing particles, when paraffin and graphite are present in the telescope: (Iparaffin Igraphite) н e-2,8/LP. In this case L denotes the range of inelastic proton-proton, interaction. The values of this ratio obtained by means of va-Card 1/2

The Cross Section of non-Elastic Proton-Proton Interaction at 20-4-13/52 Proton Energies Exceeding 7 BeV.

rious apparatus and various methods, are shown in a diagram and the corrections taken into account with this operation, are enumerated. The values $L_p^{M=51^{+23}}$ g.cm⁻¹ and of 32 1 lo mb, are obtained accorp

ding to the all data given in the table by taking the statistical weights for the range and the pross-section of the inelastic interaction proton-proton into account. In conclusion these results are still compared with those obtained by other authors.

There are 1 figure, 1 table, and 5 references, 3 of which are Slavic.

ASSOCIATION. State University imeni M. V. Lomonosov, Moscow (Moskovskiy gosudarstvennyy universitet imeni M. V. Lomonosova).

PRESENTED. June 29, 1957, by D. V. Skobel'tsyn, Academician.

SUBMITTED. June 28, 1957.

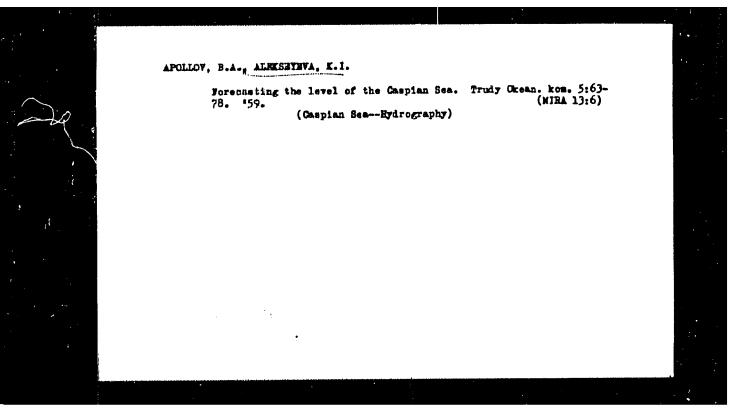
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Card 2/2

21(0) AUTHORS: Alekseyeva, K. I., Grigorov, N. L. S07/56-35-3-7/6 TITLE: Inelastic Interaction of Protons of Cosmic Radiation With Kinetic Energies Above 7 BeV With Carbon- and Hydroge Nuclei (Neuprugoye vzalmodeystviye protonov kosmicheskikh luchey a kineticheskoy energiyey boliche 7 BeV a yadrami ugleroda i vodoroda) Churnal eksperimentaliney a teoreticheskoy fiziki, 1958, Vol 35, Br 3, pp 599 - 611 (USSR) PER HOD HOAL: ABSTRACT: In the present paper the authors give a report on the results obtained by mensuring the effective cross sections of inelastic interactions of cosmic primary particles (protons) with H- and C-nuclei. (Compare also references 5-8), as well as on results obtained when determining the number of particles produced by such processes. Measurments were carried out in the stratosphere in an altatude of 20-25 km at a geomagnetic latitude of 310N. Two methods were used for determining cross sections: In one case a telescope was used which contained a lend filter as well as the graphite- and Card 1/3

Inclusive Interaction of Protons of Cosmic Rallation SOV/56-3-3-1/6* With Kisetic Energies 4bove 7 BeV With Carbon- and Hydrogen N clei

Card 2/5



SOV/56-37-2-8/56 21(7) Alekseyeva, K. I., Grigorov, N. L. AUTHORSE Heavy Nuclei Flux in Primary Cosmic Radiation at a Geomagnetic Latitude of $31\,^{\circ}\text{R}$ CIMLE: Zhurnal eksperimental'noy i teoreticheskoy fiziki, 1959, PERIODICAL: Vol 37, Nr 2(8), pp 380-388 (USSR) The flux of primary heavy particles in the stratosphere ABSTRACT: (25-27 km altitude, atmosphere: 24 g/cm-2) was measured (September 18, 1954) with an apparatus consisting of a telescope surrounded by hodoscope counters and two pulse ionization chambers placed between the trays of the telescope counters (Geiger-Müller). A scheme of this device, which transmitted its results to the earth radiographically, is shown in figure 1. An example for the recorded pulses from chamber and hodoscope is given by the reproduction of part of a film, and is discussed in detail. Figure 3 shows the spectrum of the ionization pulses of individual particles in an altitude of 1 km. A detailed chapter further deals with the evaluation of experimental data; the result is given by a table: Card 1/3

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	Heavy Nuclei	Flux in Primary Commic Radiation a	SOV/5 t a Geomagnet	6-37-2-8/56 ic Latitude of 31	N
6			Z = 2	z > 2	•
		Number of recorded individual particles (atmosphere: 24 g/cm-2)	317 ± 18	10±3.2	
		Correction with respect to a back- ground of simply charged par- ticles	-90 ±2 2		
		Correction respecting the number of nuclei producing 6-showers	+45±6		
		Number of primary heavy nuclei without nuclear interactions	272 ± 29	10±3.2	
		The same in cm 2min 1 steradian 1	0.154±0.016	0.005620.0018	
		Coefficient of the absorption of heavy nuclei due to nuclear interaction in the material of the			
	Card 2/3	device $e^{-x/L}$ (x = absorber depth, L = inelastic interaction range)	0.815	0.740	
	•	*			

SOV/56-37-2-8/56 Heavy Nuclei Flux in Primary Cosmic Radiation at a Geomagnetic Latitude of 31°S Z = 2 $\mathbf{z} > 2$ The same in the mir layer above the device 0.565 0.400 The heavy particle flux $I_{\mathbf{Z}}^{o}$ on the boundary of the atmosphere in cm⁻².min⁻¹.steradian⁻¹ 0.33520.035 0.01920.006 I_3^0/I_0 in % (I_0 = total particle flux) 16±2 In the text these data are mostly mentioned once more and discussed. V. P. Grushin took part in this work. There are 6 figures, 1 table, and 11 references, 4 of which are Soviet. ASSOCIATION: Moskovskiy gosudarstvennyy universitet (Moscow State University) SUBMITTED: March 7, 1959 Card 3/3

21(7) 507/56-37-3-2/62 AUTHORS : Alakseyeva, K. I., Brikker, S. I., Grigorov, N. L., Murzin, V. S., Skvin, F. D. TITLE: Investigation of the Production of π° -Mesons in the Stratosphere in the Case of Interaction of Protons and a Particles of Cosmic With Carbon Nuclei PERIODICAL: Zhurnal eksperimental'noy i teoreticheskoy fiziki, 1959, Vol. 37, Nr 3(9), pp 596 - 603 (USSR) ABSTRACT: In the present paper the authors describe the carrying out of and the results obtained by experiments serving the purpose of determining the average κ^0 -energy at an altitude of 25 km. The experiments themselves had taken place on September 2c, 1954 at a geomagnetic latitude of 31° N; measurement data were transmitted to the earth radiotelegraphically. For the purpose of determining \overline{E}_{RC} , the ionization in the maximum of the cancades occurring in π^0 -decay was measured. The average primary energy of the protons $E_{\rm op}$ was 20 Bev, that of the α -particles $\mathbf{E}_{\mathrm{opt}}$ amounted to 4c Bev. The experimental arrangement (schemati Card 1/4 cally shown by figure 1) consisted essentially of pulse ioni-

Investigation of the Production of π^0 -Mesons in the 30V/56-37-3-2/62 Stratosphere in the Case of Interaction of Protons and I-Particles of Cosmic Rays With Carbon Nuclei

tation chambers and a hodoscope. The counters were Geiger-Mueller counters connected in triple coincidence. The entire arrangement is described in detail. The results are shown in form of a table and by figure 2. Figure 2 in three diagrams shows the number of showers N, in which the given ionization was found, as a function of J/J_0 in the lower chamber (II). J_0 denotes the probable ionization of a relativistic simply charged particle, J the ionization of the given particle. The uppermost diagram contains the range $O((J/J_0)_1 < 30$, the middle one $3.0 < (J/J_0)_1 < 7.5$, and the third $(J/J_0) > 7.5$. The unbroken lines refer to measurements carried out with the help of a graphite filter, the dotted lines show the spectrum without such a filter. The average number of the electrons $\overline{\gamma}$ in the maximum of the γ -cascade is calculated by means of formula (2). Results:

Card 2/4

Investigation of the Production of mo-Mesons in the SOV/56-37-3-2/62 Stratosphere in the Case of Interaction of Protons and u-Particles of Cosmic Rays With Carbon Nuclei

Frimary particles: protons u-particles J/J_0 in chamber I 0 - 3.0 3.0 - 7.5 V 11.4 \pm 3.5 32.2 \pm 23.0 Number of electrons in the

avalanche N_{max} 26 ± 8 73 ± 52 $E_{\pi^0}[ev]$ (2.1±0.6)·10⁹ (5.8±4.2)·10⁹ ($E_{\pi^0}/\bar{s_0}$)·100[%] 10±3 14±10

For the determination of $R_{\rm max}$ formula (1) by Tamm and Belen'kiy is used. For the given primary energies of protons and aparticles (20 and 40 Bev respectively) the following values are obtained for the energy fraction k contributed by these particles for π^0 -production: $k_p = (10+3)\%$ and $k_z = (14+10)\%$. L. G. Landsberg participated in this work. The authors thank I. P. Ivanenke for discussions. There are 2 figures, 1 table,

Card 3/4

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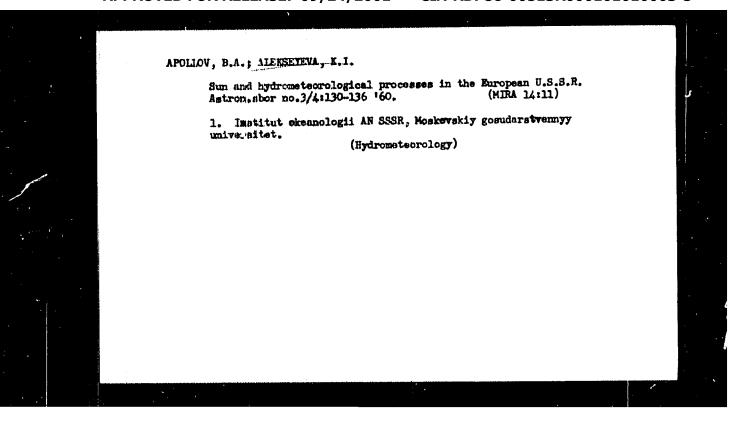
and 10 references, 7 of which are Soviet.

Investigation of the Froduction of x⁰-Mesons in the 30V/56-37-3-2/62
Stratosphere in the Case of Interaction of Protons and 3-Particles of Cosmic
Rays With Carbon Nuclei

ASSOCIATION: Moskovskiy gosudarstvennyy universitet (Moscow State University)

SUBMITTED: March 7, 1959

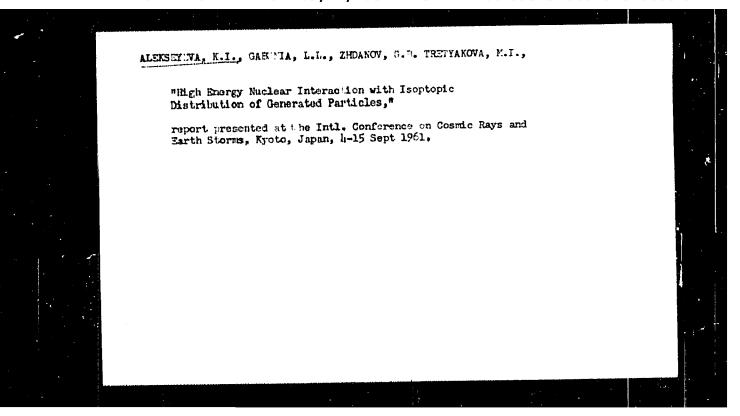
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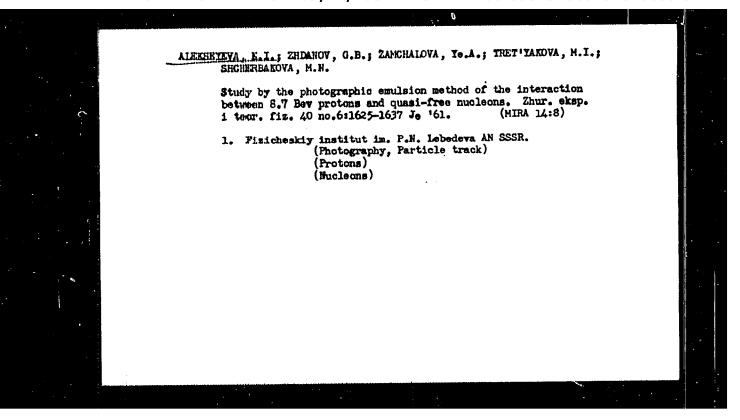


ALEXSEVEVA, K.I., GARNIYA, L.I., ZHDANOV, G.B., ZARCHALOVA, B.V.,
SHCHERLIKOVA, W.I., and TRETIAKOVA, M.I.,

"Study of Composition of Primary Cosmic Radiation at an
Altique of 320 Kilometers,"

report presented at he Intl. Conference on Cosmic R.ys and
Earth Storms, Kyoto, Japan, 1-15 Sept 1961.





ALEKSRYEVA, R. I. ZHUANOV, G. B., THETYANOVA, M. I., TEYTOVICH, V. N., and
HICHEREAROVA, M. N.

"Ionization momentum dependence for electrons in the ultra-relativistic region"

Fourth Enzernational Colloquium on Photography (Corpuscular) - Munich, West

Germany, 3-8 Sep 62

CIA-RDP86-00513R000101010003-5

3/560/62/000/012/001/014 1046/1246

AUTHORS:

Aleksews, K.I., Subuniya, L.L., Zidanov, G.B., Zanchalova, Ye.A., Shehorbistova, h.E. and Tret'yakova, K.I.

TITLE:

investigation of the primary cosmic radiation composition at an altitude of 320 km

GOUNCE:

Akademiya nauk SSSR. Iskusotvennyye sputniki Zemli, no. 12, Mosess, 1962, 6-15

TEXT: The automatic apparatus whose design was reported at the International Conference on Nuclear Photography (1960) is applied to impulse and ionization measurements of middle-weight coarie nuclei. In sultiple scattering measurements, the time required to measure one 10 mm trail in 7 minutes; in ionization measurements, 30 minutes per trail are required. This is at least 5 times an fact as in visual measurements. The resolution of the apparatus in ordinary direcumtances is sufficient to separate between the Li, Be, B and C, N, O groups. Instrumental errors, however, reduce the accuracy of measuring trail discontinuities by up to 30-40. As compared with visual measurements for a given

Card 1/2

